# Orange Book: Approved Drug Products with Therapeutic Equivalence Evaluations

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**Search Results for Proprietary Name, Active Ingredient or Application Number: *021636***

RX OTC DISCN

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| Mkt. Status | Active Ingredient | Proprietary Name | Appl. No. | Dosage Form | Route | Strength | TE Code | RLD | RS | Applicant Holder |
|---|---|---|---|---|---|---|---|---|---|---|
| DISCN | OMEPRAZOLE; SODIUM BICARBONATE | ZEGERID | N021636 | FOR SUSPENSION | ORAL | 20MG/PACKET; 1.68GM/PACKET | | RLD | | SALIX PHARMACEUTICALS INC |
| DISCN | OMEPRAZOLE; SODIUM BICARBONATE | ZEGERID | N021636 | FOR SUSPENSION | ORAL | 40MG/PACKET; 1.68GM/PACKET | | RLD | | SALIX PHARMACEUTICALS INC |
| **Mkt. Status** | **Active Ingredient** | **Proprietary Name** | **Appl. No.** | **Dosage Form** | **Route** | **Strength** | **TE Code** | **RLD** | **RS** | **Applicant Holder** |

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**Product Details for NDA 021636**

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ZEGERID (OMEPRAZOLE; SODIUM BICARBONATE)
20MG/PACKET;1.68GM/PACKET
Marketing Status: Discontinued

**Active Ingredient:** OMEPRAZOLE; SODIUM BICARBONATE
**Proprietary Name:** ZEGERID
**Dosage Form; Route of Administration:** FOR SUSPENSION; ORAL
**Strength:** 20MG/PACKET;1.68GM/PACKET
**Reference Listed Drug:** Yes
**Reference Standard:** No
**TE Code:**
**Application Number:** N021636
**Product Number:** 001
**Approval Date:** Jun 15, 2004
**Applicant Holder Full Name:** SALIX PHARMACEUTICALS INC
**Marketing Status:** Discontinued
Patent and Exclusivity Information

ZEGERID (OMEPRAZOLE; SODIUM BICARBONATE)
40MG/PACKET;1.68GM/PACKET
Marketing Status: Discontinued

**Active Ingredient:** OMEPRAZOLE; SODIUM BICARBONATE
**Proprietary Name:** ZEGERID
**Dosage Form; Route of Administration:** FOR SUSPENSION; ORAL
**Strength:** 40MG/PACKET;1.68GM/PACKET
**Reference Listed Drug:** Yes
**Reference Standard:** No
**TE Code:**
**Application Number:** N021636
**Product Number:** 002
**Approval Date:** Dec 21, 2004
**Applicant Holder Full Name:** SALIX PHARMACEUTICALS INC
**Marketing Status:** Discontinued
Patent and Exclusivity Information